GONIOTAIRE- hypromellose 2906 (4000 mpa.s) solution
Altaire Pharmaceuticals Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Goniotaire

Altaire

Goniotaire

Hypromellose 2.5% Ophthalmic

Demulcent Solution

15mL

NDC 59390-182-13

Drug Facts

Each mL Contains:

Active:

Hypromellose 25mg (2.5%):

Inactives:

Benzalkonium Chloride, Boric Acid, Edetate Disodium, Sodium Borate, Water for injection. Hydrochloric Acid and/or sodium hydroxide may be added to adjust pH (6.0 to 7.8).

NOTE: If this solution dries on optical surfaces, let them stand in cool water before cleansing.

INDICATIONS:

For professional use in gonioscopic examinations.

DIRECTIONS:

Fill gonioscopic prism with solution as necessary.

WARNINGS:

To avoid contamination do not touch tip of container to any surface. Replace cap after using. Not for use in conjunction with hot laser treatment. If solution changes color or

becomes cloudy, do not use.

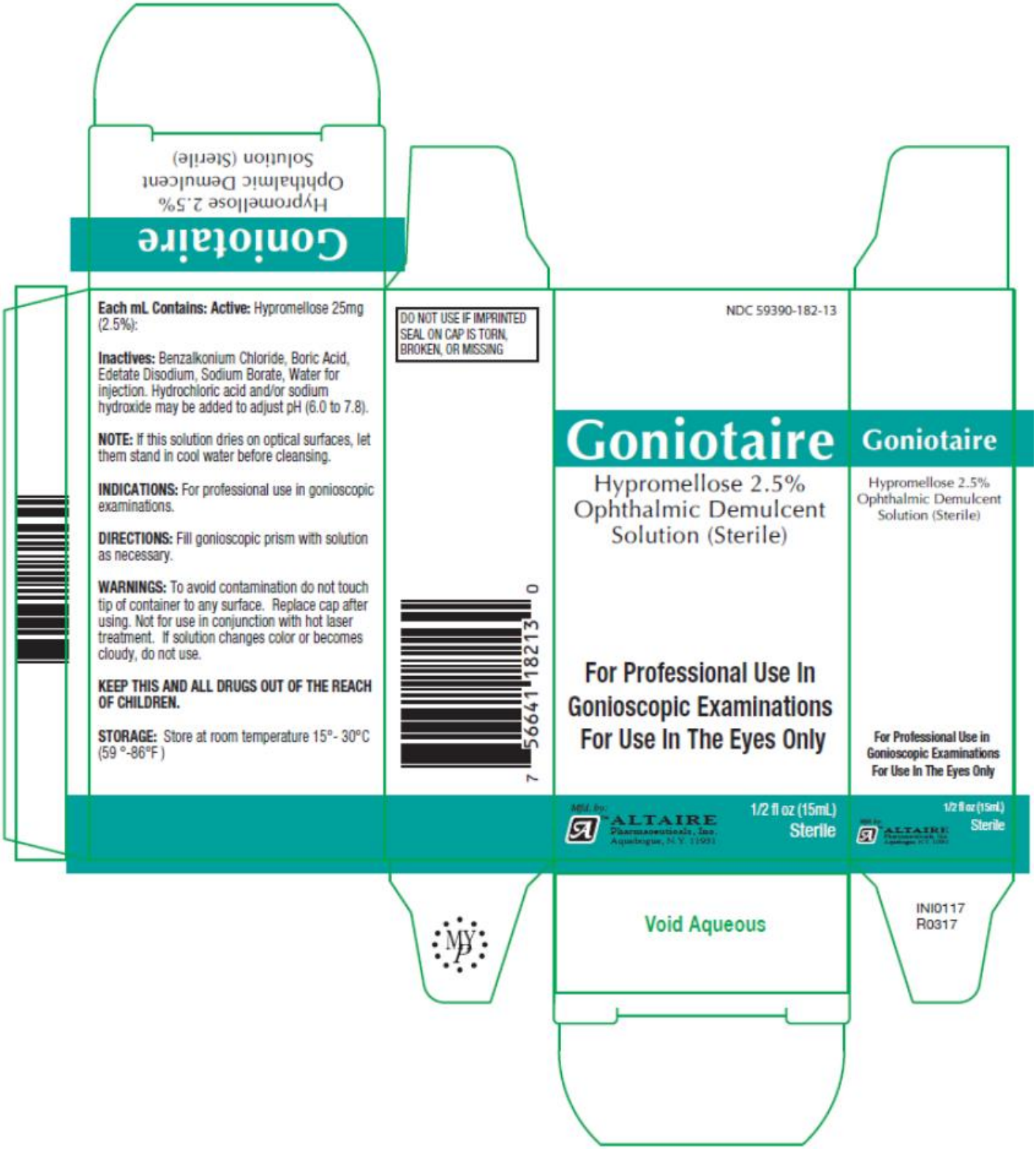
KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

STORAGE:

Store at room temperature 15°- 30°C (59°- 86°F).

PRINCIPAL DISPLAY PANEL

NDC 59390-182-13
Goniotaire
Hypromellose 2.5%
Ophthalmic Demulcent
Solution (Sterile)
½ fl oz (15mL)
sterile



GONIOTAIRE

hypromellose 2906 (4000 mpa.s) solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:59390-182
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYPROMELLOSE 2906 (4000 MPA.S) (UNII: 5EYA69XGAT) (HYPROMELLOSE 2906 (4000 MPA.S) - UNII:5EYA69XGAT)	HYPROMELLOSE 2906 (4000 MPA.S)	25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	**HYDROCHLORIC ACID** (UNII: QTT17582CB)	
May contain	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59390-182-13	1 in 1 CARTON	01/18/2002	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/18/2002	

Labeler - Altaire Pharmaceuticals Inc. (786790378)

Revised: 10/2022 Altaire Pharmaceuticals Inc.